Social-bond strength influences vocally-mediated recruitment to mobbing

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Strong social bonds form between individuals in many group-living species, and these relationships can have important fitness benefits. When responding to vocalisations produced by groupmates, receivers are expected to adjust their behaviour depending on the nature of the bond they share with the signaller. Here we investigate whether the strength of the signaller–receiver social bond affects response to calls that attract others to help mob a predator. Using field-based playback experiments on a habituated population of wild dwarf mongooses (*Helogale parvula*), we first demonstrate that a particular vocalisation given on detecting predatory snakes does act as a recruitment call; receivers were more likely to look, approach and engage in mobbing behaviour than in response to control close calls. We then show that individuals respond more strongly to these recruitment calls if they are from groupmates with whom they are more strongly bonded (those with whom they preferentially groom and forage). Our study therefore provides novel evidence about the anti-predator benefits of close bonds within social groups.

Introduction

A common feature of stable social groups is the presence of close bonds, or friendships', between individuals [1,2]. While there are many different ways to quantify the strength of such relationships [3], it is recognised that 'strong' bonds with groupmates can provide considerable long-term health and fitness benefits [1,2]. However, less is known about potential short-term survival benefits [1,4]. Reduction of predation risk is facilitated in many species by a range of different acoustic signals that can induce fleeing, increase vigilance and coordinate defensive actions [5,6]. Recent work on chimpanzees (*Pan troglodytes*) and yellow-bellied marmots (*Marmota flaviventris*) has shown that the propensity of individuals to *give* flee alarm calls can

depend on the presence of close affiliates and their own position in a social network [7,8]. Behavioural adjustments in *response* to at least some anti-predator vocalisations (e.g. those that coordinate defence) might also be expected depending on the level of affiliation with the caller, but little attention has been paid to receivers in this regard (see [4] for an exception).

In many taxa, certain vocalisations serve to attract others to the caller. These 'recruitment' calls often advertise the location of a food source [9], but are also given when individuals encounter specific predators [10]. Predator-related recruitment calls can engage both conspecifics and heterospecifics in collective mobbing behaviour, with responders purposely approaching and harassing the threat [10–12]. Mobbing is costly in terms of potential injury or death, lost foraging time, and the risk of attracting further predators [13–15]. Like many other vocalisations, predator-related recruitment calls can convey information about the caller's identity [4,16]. However, only one empirical study has considered how within-group signaller–receiver bond strength might influence call responses: crested macaques (*Macaca nigra*) oriented for longer towards a loudspeaker playing recruitment calls of close affiliates compared to those of weak affiliates [4].

Here we use field playback experiments to examine whether caller identity influences receiver responses to the calls given by dwarf mongooses (*Helogale parvula*) on encountering predatory snakes. Having first demonstrated that these calls do indeed function to recruit group members, we investigate the role of social-bond strength between callers and responders. Specifically, we test whether individuals show greater responses to the recruitment calls of individuals to which they are more strongly bonded.

2. Material and Methods

(a) Study site and population

Data were collected on Sorabi Rock Lodge Reserve, South Africa from nine wild dwarf mongoose groups habituated to close observation [17,18]; full methodology in Supplementary Material (SM); datasets available in [19]. Data on natural mobbing events – approaching, cooperative harassing and attacking of a predator – were collected using all-occurrence sampling between January 2014 and March 2016.

(b) Playback experiment 1

To test whether the calls given by dwarf mongooses when they detect a predator to be mobbed (see Results) function to recruit others, we compared responses to playback of these calls and control close calls given while foraging (Fig. SM1). Putative 'recruitment' calls were recorded during natural snake-mobbing events and rubber-snake presentations. Close calls were recorded opportunistically during foraging bouts. Nine randomly selected subordinate individuals received separate 10-min playbacks of the two call types at natural rates and amplitudes. Playbacks to the same focal individual were of calls from the same adult subordinate group member and were separated by 1 h; the presentation order of the two playback types was alternated to different focal individuals. Focal individuals were filmed during playback, and data on looking, approaching and mobbing behaviour subsequently extracted.

(c) Playback experiment 2

To assess how the response to recruitment calls is influenced by signaller–receiver social-bond strength, we conducted a second playback experiment. Eight individuals from four groups (those with sufficient subordinate group members to enable comparison of a stronger and weaker social bond) each received two 10-min playbacks of recruitment calls, one from a subordinate groupmate with whom they shared a relatively strong bond and one with whom they shared a relatively weak bond. Social-bond strengths were determined from composite sociality indexes (CSI) [4,20] based on grooming and nearest-neighbour foraging distances. The use of multiple behavioural indices strengthens the assessment of bond strength, and previous research has established that grooming and foraging associations are strongly correlated within dwarf mongoose groups (full details in SM). Experimental signaller–receiver dyads were selected to maximise the difference in CSI scores for a given focal individual. Playbacks to the same focal individual were separated by 7.5±2.3 days (mean±SE; range: 2–15); group size was the same for both trials to the same individual. Variation in the time between trials to the same focal individual did not significantly affect either the absolute response shown in the second trial (Jonckheere-Terpstra test, duration of looking: T_{JT}=17, N=8, P=0.24; duration of physical response: T_{JT}=11, N=8, P=0.61) or the difference in response between the two trials (duration of looking: T_{JT}=12, N=8, P=0.90; duration of physical response: T_{JT}=15, N=8, P=0.43). The presentation order of the two playbacks was

alternated to different focal individuals. Focal individuals were filmed, and data extracted, as in Experiment 1.

(d) Statistical analysis

The response of focal foragers to the two types of call (Experiment 1) were analysed using two McNemar related-samples tests (for tendencies to look at and to approach the speaker) and two Wilcoxon signed-rank tests (for durations of looking and physical responses; the latter defined as the time spent approaching and mobbing). Data from Experiment 2 were analysed using linear mixed models (LMMs) and generalized linear mixed models (GLMMs), to account for data collection from more than one focal individual per group. For all models, the fixed effects of social-bond strength (strong, weak), group size and trial order (1, 2) were fitted, and focal individual nested in group was included as a random term.

3. Results

Sixty-one natural mobbing events were observed in response to snakes (puff adders (*Bitis arietans*), Mozambique spitting cobras (*Naja mossambica*), black mambas (*Dendroaspis polylepis*), African rock pythons (*Python sebae*)). In all cases, the first individual to locate the threat gave a particular vocalisation (Fig. SM1a); this was the vocalisation tested in the playback experiments. Other group members approached the caller, searched for the threat and then surrounded the predator, displaying typical mobbing behaviours such as head bobbing and weaving, striking at the predator, and threat scratching. Mobbing events lasted for 697 ± 148 s (mean ± SE) and involved 62% ± 4% of the group.

Compared to close-call playback, playback of calls given on detecting snakes (see above) resulted in focal foragers being more likely to look at the speaker (McNemar's test: N=9 paired playbacks, P=0.013), looking for longer (Wilcoxon signed-rank test: Z=0, N=9, P=0.004), being more likely to approach the speaker (McNemar's test: N=9 paired playbacks, P=0.041) and responding physically for longer (Wilcoxon signed-rank test: Z=0, N=9, P=0.014).

Controlling for a significant negative effect of trial order in several cases (Table SM1), focal foragers were more likely to look at the speaker (GLMM: χ^2=4.56, df=1, P=0.033; Fig. 1a), looked for longer (LMM: χ^2=11.06, df=1, P=0.001; Fig. 1b), were more likely to approach the speaker (GLMM: χ^2=10.62, df=1, P=0.001; Fig. 1c), and responded physically for longer (LMM: χ^2=854.95, df=1, P<0.001; Fig. 1d) when played recruitment calls from individuals to

which they were strongly bonded compared to those from groupmates to which they were more weakly bonded.

4. Discussion

Our study shows that, on detecting predatory snakes, dwarf mongooses produce specific vocalisations that act as recruitment calls. These calls increase the likelihood of the caller being joined by other group members in mobbing the threat, as is the case in various other species [8,9]. We demonstrate experimentally that the response to these recruitment calls differs depending on the social-bond strength shared by the signaller and receiver. Individuals showed a greater response (in terms of looking, approaching and mobbing) when hearing recruitment calls from groupmates to which they were strongly bonded compared to those with which they shared a weaker bond. Although a previous study indicated that crested macaques orientated more to (i.e. looked in the direction of) the recruitment calls of close affiliates than weak affiliates, they found no difference in the tendency to approach or duration of response [3]. To our knowledge, the current work is therefore the first to show greater active responses to the recruitment calls of groupmates with whom receivers share stronger bonds (see [21] for an example of how long-term familiarity increases the likelihood that neighbours assist one another in nest defence).

Heightened responses to the recruitment calling of particular group members could theoretically be a by-product of factors influencing the formation of social bonds. If individuals were more likely to form strong bonds with groupmates of similar age and size, for example, dyads with strong bonds would have similar risk profiles. Mobbing behaviour by one of these other individuals would thus be a potentially good indication of a threat to self. Within dwarf mongoose groups, however, there is much variation in social-bond strength between individuals of the same age (JM Kern unpub. data). Indeed, in several cases, the strongly and weakly bonded experimental individuals were littermates. Instead, the preferential response to recruitment calls from strongly bonded groupmates may arise from a trade-off between the benefits and costs, given that mobbing behaviour is costly [11–13]. There are a number of potential such possibilities.

First, it has been suggested that mobbing may function as a costly signal, advertising individual quality to conspecifics [17]. Individuals may invest more in signalling their quality to those

with which they share strong bonds to uphold their attractiveness as a close partner, though so far support for this hypothesis is lacking [10,18]. Second, individuals may preferentially associate with close affiliates in stressful situations. In pilot whales (*Globicephala melas*), for example, closely affiliated dyads increase their synchronization when swimming in stressful circumstances [19]. Third, there may be variation in the relative costs and benefits of responding to callers with whom receivers have stronger or weaker bonds. The effectiveness of mobbing increases with the number of participants [13], thus groupmates may directly improve the survival chances of a caller when they respond to recruitment calls. Reciprocal cooperation, often performed over long time periods, may also be more likely between strongly bonded individuals [20]. Receivers who respond to close affiliates now may therefore stand to gain future advantages, including likely assistance themselves in future mobbing events or intra-group conflicts [21], in addition to the ongoing advantages of close friendships.

Recent experimental work using other call types has demonstrated an effect of social-bond strength and other social attributes on caller behaviour [7, 8]. Here, we show an effect of social bonds on receiver responses (see also [3]), enhancing our understanding of the role of social bonds in intra-group interactions. While the long-term benefits of close social bonds are well established, particularly in primates, the potential in other species and in the context of predation has been little explored. In general, by adjusting their responses depending on caller identity, receivers can facilitate more efficient and effective use of social information.

Ethics This study was conducted under all required ethical approvals.

Data accessibility All data for this paper will be archived in Dryad.

Author contributions J.M.K. & A.N.R. designed the study; J.M.K. collected the data; J.M.K. analysed the data with advice from A.N.R.; J.M.K & A.N.R. interpreted the data and co-wrote the paper.

Competing interests We have no competing interests.

Funding The work was supported by a University of Bristol studentship to J.M.K.

Acknowledgements We thank B. Rouwhorst and H. Yeates for access to their land, C. Esterhuizen for logistical support and 12 research assistants for observational data collection.

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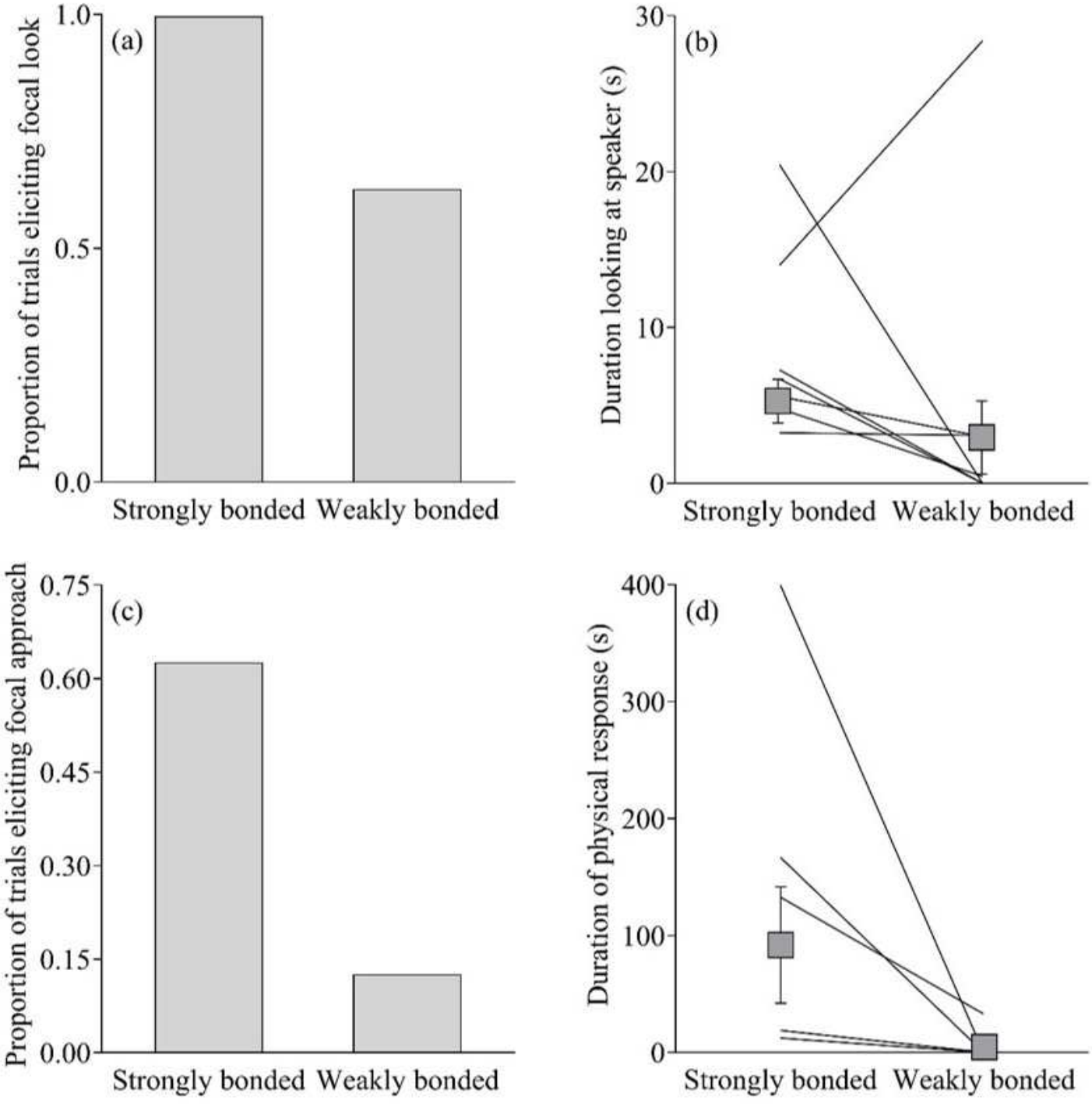
Figure Legends

Figure 1 Response of dwarf mongooses to the playback of recruitment calls given by groupmates to which they are strongly or weakly bonded. (a) Proportion of trials eliciting looking at speaker, (b) total duration looking at speaker, (c) proportion of trials eliciting approach to speaker, and (d) total duration of physical response. For (a)–(c), *N*=eight individuals, four groups; for (d), *N*=seven individuals, three groups. Shown for (b) and (d) are results for each focal individual separately (lines) and the overall treatment mean (solid squares) ± SE.

Figure 1


(a)
Proportion of trials eliciting focal look
1.0
0.5
0.0
Strongly bonded
Weakly bonded
(b)
Duration looking at speaker (s)
30
20
10
0
Strongly bonded
Weakly bonded
(c)
Proportion of trials eliciting focal approach
0.75
0.60
0.45
0.30
0.15
0.00
Strongly bonded
Weakly bonded
(d)
Duration of physical response (s)
400
300
200
100
0
Strongly bonded
Weakly bonded